

## Health Update:

### Rabies Manufacturer's Recall of Human Rabies Vaccine

April 9, 2004

This document will be updated as new information becomes available. The current version can always be viewed at <http://www.dhss.state.mo.us/>.

The Missouri Department of Health & Senior Services (DHSS) is now using 4 types of documents to provide important information to medical and public health professionals, and to other interested persons:

**Health Alerts** convey information of the highest level of importance which warrants immediate action or attention from Missouri health providers, emergency responders, public health agencies, and/or the public.

**Health Advisories** provide important information for a specific incident or situation, including that impacting neighboring states; may not require immediate action.

**Health Guidances** contain comprehensive information pertaining to a particular disease or condition, and include recommendations, guidelines, etc. endorsed by DHSS.

**Health Updates** provide new or updated information on an incident or situation; can also provide information to update a previously sent Health Alert, Health Advisory, or Health Guidance; unlikely to require immediate action.

Health Update  
April 9, 2004

**FROM: RICHARD C. DUNN  
DIRECTOR**

**SUBJECT: Manufacturer's Recall of Human Rabies Vaccine**

[This **Health Update** provides additional information on the manufacturer's recall of human rabies vaccine **Health Advisory issued on April 3, 2004.**)

The attached document is an algorithm for triage and disposition of persons who received Imovax rabies vaccine from 9/23/03 through 4/2/04 from vaccine lots X0667-2, X0667-3, W1419-2, and W1419-3. It was developed jointly by the New York State Health Department and the Centers for Disease Control and Prevention.

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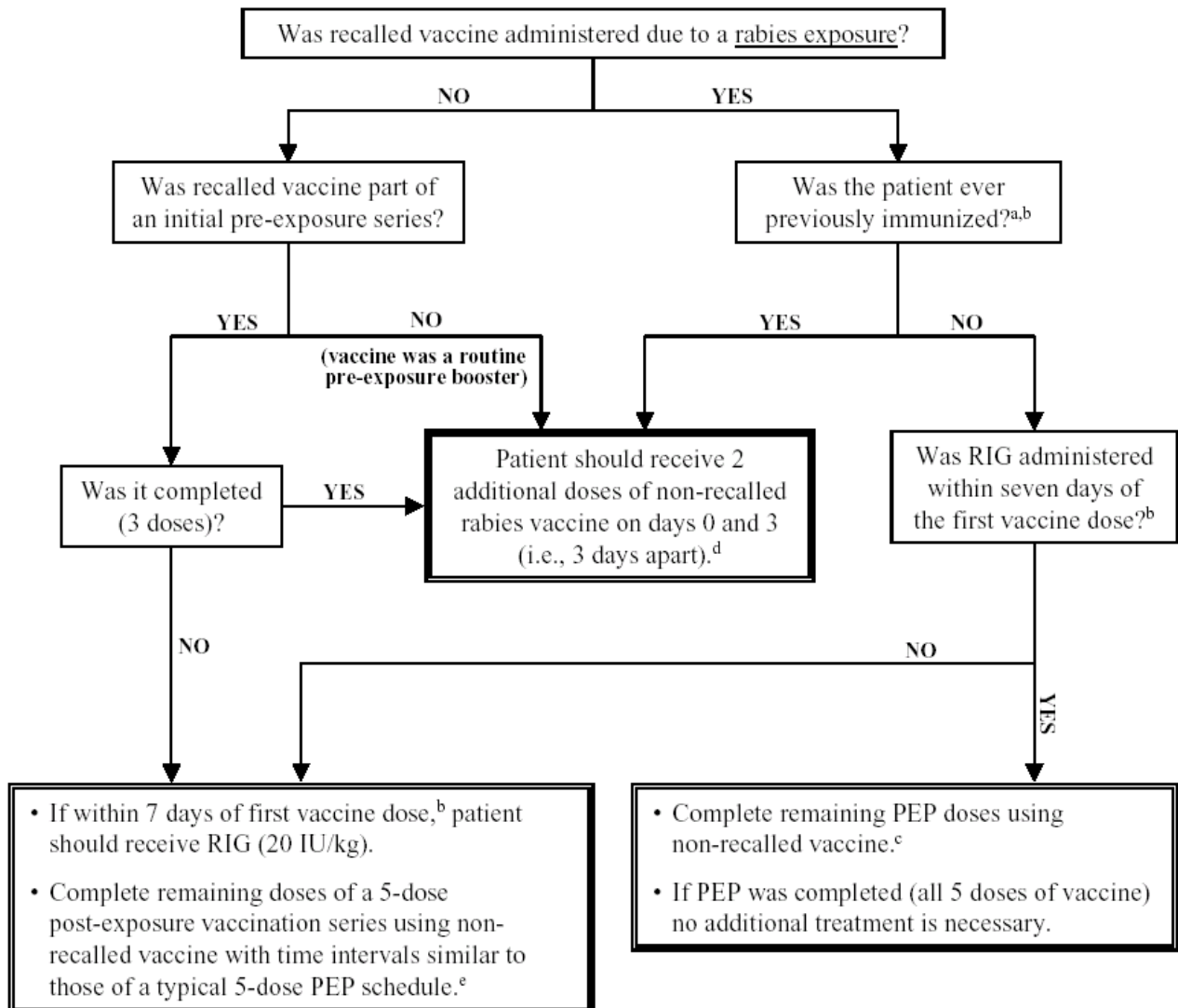
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**Questions should be directed to the Department of Health and Senior Services  
24 hours a day, 7 days a week at 800-392-0272.**

## Patient\* Triage, Imovax Rabies Vaccine Recall, April 2, 2004

(\*who received Imovax rabies vaccine from 9/23/03 through 4/2/04  
from vaccine lots X0667-2, X0667-3, W1419-2, and W1419-3)



<sup>a</sup> Previously immunized = 3 previous doses for pre-exposure or post-exposure.

<sup>b</sup> Any vaccine regardless of whether that vaccine was in the recalled lots.

<sup>c</sup> If PEP was discontinued because the animal was found to not have rabies, the patient should complete series using non-recalled vaccine (i.e., total of 5 doses of vaccine including any already given.)

<sup>d</sup> If < 7 days since completion of 3 doses, the additional 2 doses should be given at days 7 and 21 after last previous dose.

<sup>e</sup> Typical 5-dose PEP schedule = vaccine on days 0, 3, 7, 14, 28.